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IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Application of:

BLECHA, FRANK, et al.

Serial No. : 10/014,147

Filed: December 07, 2001

SYNTHETIC PEPTIDES THAT INHIBIT
LEUKOCYTE SUPEROXIDE ANION
PRODUCTION AND/OR ATTRACT
LEUKOCYTES

Docket No. 23625-DIV1

Group Art Unit No. 1653

Examiner:

U.S. Patent and Trademark Office
Arlington, VA 22202

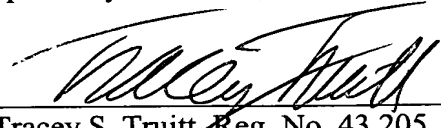
Sir:

SEQUENCE TRANSMITTAL

Transmitted herewith are: Express Mail Transmittal letter (1 page); Sequence Transmittal letter (1); 2 CFR CD's Compact Discs (2 compact disks); Sequence Listing (5) and Copy of Notice to Comply with Requirement for Patent Applications Containing Nucleotide Sequence and/or Amino Acid Sequence Disclosures (2 pages); CRF Sequence Statement Transmittal Letter (1 page); and a return postcard. These documents are being deposited with the United States Postal Service "Express Mail Post Office to Addressee" service under 37 CFR 1.10 in an envelope addressed to: U.S. Patent and Trademark Office; Box Sequence; P.O. Box 2327; Arlington, VA 22202 on October 2, 2002.

EV 110672189 US

Respectfully submitted,

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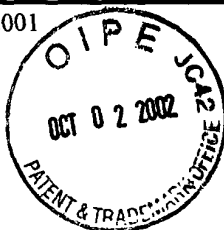
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APPLICATION NUMBER	FILING/RECEIPT DATE	FIRST NAMED APPLICANT	ATTORNEY DOCKET NUMBER
10/014,147	12/07/2001	Frank Blecha	23625-DIV1

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CONFIRMATION NO. 7389

FORMALITIES LETTER



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Date Mailed: 09/23/2002

**NOTICE TO COMPLY WITH REQUIREMENTS FOR PATENT APPLICATIONS
CONTAINING NUCLEOTIDE SEQUENCE AND/OR AMINO ACID SEQUENCE
DISCLOSURES**

Filing Date Granted

Applicant is given **TWO MONTHS FROM THE DATE OF THIS NOTICE** within which to file the items indicated below to avoid abandonment. Extensions of time may be obtained under the provisions of 37 CFR 1.136(a).

- A copy of the "Sequence Listing" in computer readable form has not been submitted as required by 37 C.F.R. 1.821(e). If the effective filing date is on or after September 8, 2000, see the final rulemaking notice published in the Federal Register at 65 FR 54604 (September 8, 2000) and 1238 OG 145 (September 19, 2000). Applicant must provide an initial computer readable form (CRF) copy of the "Sequence Listing" and a statement that the content of the sequence listing information recorded in computer readable form is identical to the written (on paper or compact disc) sequence listing and, where applicable, includes no new matter, as required by 37 CFR 1.821(e), 1.821(f), 1.821(g), 1.825(b), or 1.825(d). If applicant desires the sequence listing in the instant application to be identical with that of another application on file in the U.S. Patent and Trademark Office, such request in accordance with 37 CFR 1.821(e) may be submitted in lieu of a new CRF.

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